Refine Search

Search Results -

Terms	Documents
L7 and crystalline	3

Database:

US Pre-Grant Publication Full-Text Database
US Patents Full-Text Database
US OCR Full-Text Database
EPO Abstracts Database
JPO Abstracts Database
Derwent World Patents Index
IBM Technical Disclosure Bulletins

Search: L8

Refine Search

Recall Text | Clear | Interrupt

Search History

DATE: Wednesday, November 30, 2005 Printable Copy Create Case

Set Name side by side	Query	Hit Count	Set Name result set
DB=USPT,USOC,EPAB,JPAB,DWPI,TDBD; PLUR=YES; OP=ADJ			
L8	L7 and crystalline	3	L8
L7	l5 and (562/$ or 514/$)	24	L7
L6	L5 and (polymorph$6 or orthorhombic$5)	0	L6
L5	phenoxyalkyl carboxy$8	57	L5
L4	PHENOXYALKYLCARBOXYL	0	L4
DB=USPT; PLUR=YES; OP=ADJ			
L3	4985585	2	L3
L2	4985585.pn.	1	L2
DB=PGPB; PLUR=YES; OP=ADJ			
L1	20040267041	1	L1

END OF SEARCH HISTORY

Hit List

First Hit | Clear | Generate Collection | Print | Fwd Refs | Bkwd Refs | Generate OACS

Search Results - Record(s) 1 through 3 of 3 returned.

☐ 1. Document ID: US 5707989 A

Using default format because multiple data bases are involved.

L8: Entry 1 of 3 File: USPT Jan 13, 1998

US-PAT-NO: 5707989
DOCUMENT-IDENTIFIER: US 5707989 A

TITLE: Pyrimido[5,4-D]pyrimidines, medicaments comprising these compounds, their use and processes for their preparation

DATE-ISSUED: January 13, 1998

INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Himmelsbach; Frank	Mittelbiberach			DE
von Ruden; Thomas	Wien			AT
Dahmann; Georg	Biberach			DE
Metz; Thomas	Wien			AT

US-CL-CURRENT: 514/228.2; 514/183, 514/234.2, 514/262.1, 514/81, 544/122, 544/61

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments | Claims | KWIC | Draw. De

☐ 2. Document ID: US 5177106 A

L8: Entry 2 of 3 File: USPT Jan 5, 1993

US-PAT-NO: 5177106
DOCUMENT-IDENTIFIER: US 5177106 A

TITLE: 4-amino substituted phenoxyalkyl carboxylic acid, ester, and alcohol derivatives as antihypercholesterolemic and antiatherosclerotic agents

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments | Claims | KWIC | Draw. De

☐ 3. Document ID: US 5166398 A

L8: Entry 3 of 3 File: USPT Nov 24, 1992

US-PAT-NO: 5166398
DOCUMENT-IDENTIFIER: US 5166398 A

TITLE: 4-oxy-substituted phenoxyalkyl carboxylic acid, ester, and alcohol derivatives as antihyper-cholesterolemic and antiatherosclerotic agents

Terms	Documents
L7 and crystalline	3

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     (FILE 'HOME' ENTERED AT 10:28:49 ON 30 NOV 2005)

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                  S L1

     FILE 'REGISTRY' ENTERED AT 10:29:50 ON 30 NOV 2005
L2              10 S L1 FULL

     FILE 'CAPLUS' ENTERED AT 10:29:53 ON 30 NOV 2005
L3              12 S L2 FULL
L4               7 S L3 AND PY<2003
L5               0 S L4 AND ORTHORHOMBIC
L6               0 S L4 AND POLYMORPH?

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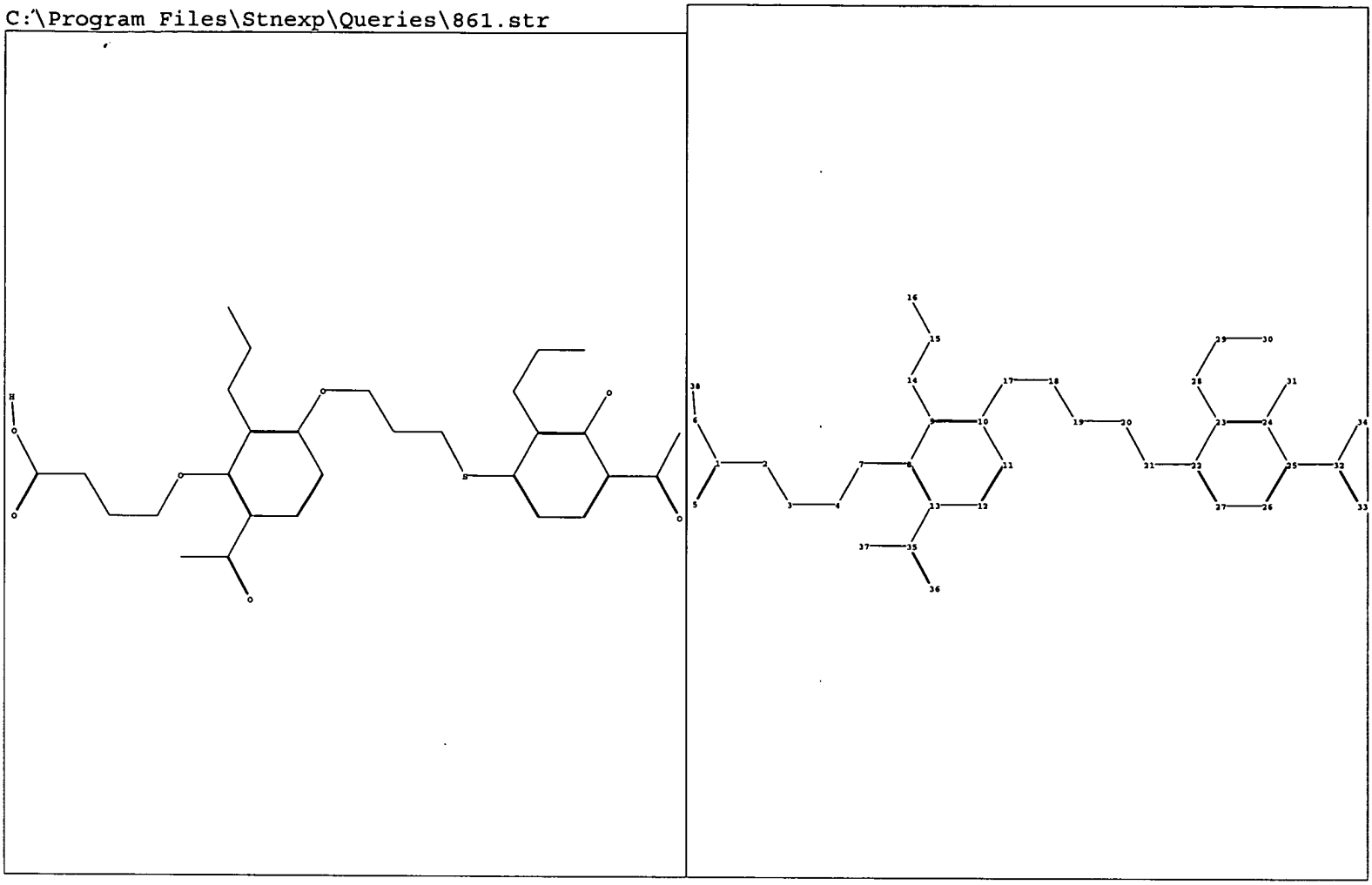

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ring nodes :
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chain bonds :
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ring/chain bonds :
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ring bonds :
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exact/norm bonds :
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35-37
exact bonds :
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normalized bonds :
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Match level :
1:CLASS 2:CLASS 3:CLASS 4:CLASS 5:CLASS 6:CLASS 7:CLASS 8:Atom 9:Atom 10:Atom
11:Atom 12:Atom 13:Atom 14:CLASS 15:CLASS 16:CLASS 17:CLASS 18:CLASS 19:CLASS
20:CLASS 21:CLASS 22:Atom 23:Atom 24:Atom 25:Atom 26:Atom 27:Atom 28:CLASS
29:CLASS 30:CLASS 31:CLASS 32:CLASS 33:CLASS 34:CLASS 35:CLASS 36:CLASS 37:CLASS
38:CLASS

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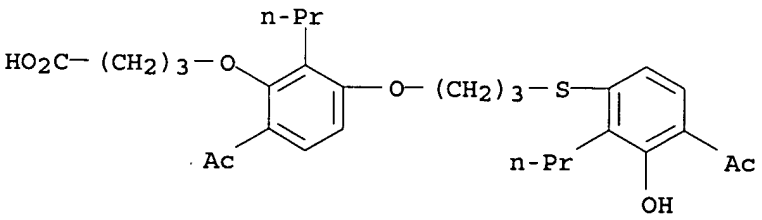
L4 ANSWER 1 OF 7 CAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2001:564826 CAPLUS
DOCUMENT NUMBER: 135:142249
TITLE: Eye drop compositions containing leukotriene antagonist KCA-757
INVENTOR(S): Kodaira, Hiromichi; Kozuka, Hitoshi
PATENT ASSIGNEE(S): Kyorin Pharmaceutical Co., Ltd., Japan
SOURCE: PCT Int. Appl., 17 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: Japanese
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001054684	A1	20010802	WO 2001-JP430	20010124 <--
W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
CA 2397755	AA	20010802	CA 2001-2397755	20010124 <--
AU 2001028804	A5	20010807	AU 2001-28804	20010124 <--
EP 1250924	A1	20021023	EP 2001-946788	20010124 <--
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
TW 526061	B	20030401	TW 2001-90101616	20010129
US 2003083378	A1	20030501	US 2002-181436	20020725
PRIORITY APPLN. INFO.:			JP 2000-17403 A	20000126
			WO 2001-JP430 W	20010124

AB Disclosed are eye drops containing a potent and selective leukotriene antagonist. Specifically, stable eye drops of an aqueous solution or suspension type, containing as the active ingredient 4-[6-acetyl-3-[3-[(4-acetyl-3-hydroxy-2-propylphenyl)thio]-propoxy]-2-propylphenoxy]butyric acid (KCA-757). An eye drop composition containing KCA-757 0.5 g, 0.1 M NaOH 20 mL, potassium dihydrogenphosphate 0.004, sodium hydrogenphosphate 0.089, NaCl 0.8 g, and 0.1 M HCl q.s. to pH 8.5, and water q.s. to 100 mL was formulated.

IT **125961-82-2**
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(eye drop compns. containing leukotriene antagonist KCA-757)

RN 125961-82-2 CAPLUS
CN Butanoic acid, 4-[6-acetyl-3-[3-[(4-acetyl-3-hydroxy-2-propylphenyl)thio]propoxy]-2-propylphenoxy]- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 25 THERE ARE 25 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 2 OF 7 CAPLUS COPYRIGHT 2005 ACS on STN

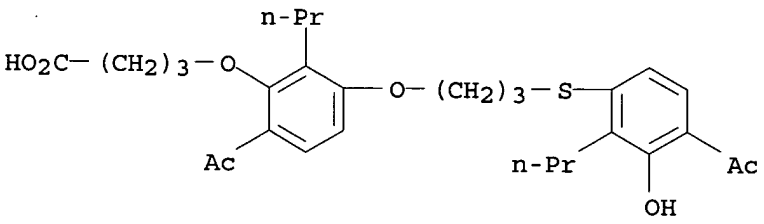
ACCESSION NUMBER: 1999:205557 CAPLUS
DOCUMENT NUMBER: 130:287054
TITLE: Powder inhalants containing [(propylphenyl)thio]propoxy]propylphenoxybutyrate for the treatment of asthma
INVENTOR(S): Hoshino, Ryoichi
PATENT ASSIGNEE(S): Kyorin Pharmaceutical Co., Ltd., Japan
SOURCE: Jpn. Kokai Tokkyo Koho, 6 pp.
CODEN: JKXXAF
DOCUMENT TYPE: Patent
LANGUAGE: Japanese
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 11079985	A2	19990323	JP 1997-251280	19970901 <--
PRIORITY APPLN. INFO.:			JP 1997-251280	19970901

AB Powder inhalants for the treatment of asthma comprise powdery 4-[6-acetyl-3-[3-(4-acetyl-3-hydroxy-2-propylphenylthio)propoxy]-2-propylphenoxy]butyric acid (I) as an active ingredient. I in combination with lubricants is suspended in an aqueous solution of polymers and spray dried to give a fine powder having an average particle diam ≤6 μm. The powders show little self-cohesive properties and little adhesion to a dispersing device. Hydroxypropyl Me cellulose 1.5 g was dissolved in 380 g distilled water and to the solution 0.5 g sucrose fatty acid ester was added, followed by 18 g I. The dispersion was subjected to a high-pressure homogenization and spray-drying to give a dry powder inhalant.

IT **125961-82-2**
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(manufacture of antiasthmatic powder inhalants containing [(propylphenyl)thiopropoxy]propylphenoxybutyrate and polymers and lubricants)

RN 125961-82-2 CAPLUS
CN Butanoic acid, 4-[6-acetyl-3-[3-[(4-acetyl-3-hydroxy-2-propylphenyl)thio]propoxy]-2-propylphenoxy]- (9CI) (CA INDEX NAME)



L4 ANSWER 3 OF 7 CAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 1996:379374 CAPLUS
DOCUMENT NUMBER: 125:58104
TITLE: Preparation of phenoxycarboxylic acid derivatives as antiallergy agents
INVENTOR(S): Matsumoto, Toyomi; Ishiguro, Juji; Myashita, Kunio; Kitamura, Genichi
PATENT ASSIGNEE(S): Kyorin Seiyaku Kk, Japan
SOURCE: Jpn. Kokai Tokkyo Koho, 4 pp.
CODEN: JKXXAF
DOCUMENT TYPE: Patent
LANGUAGE: Japanese
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 08081412	A2	19960326	JP 1994-244636	19940913 <--
PRIORITY APPLN. INFO.:			JP 1994-244636	19940913
OTHER SOURCE(S):	CASREACT 125:58104; MARPAT 125:58104			

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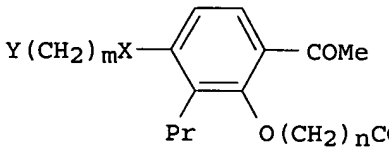

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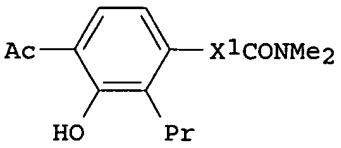

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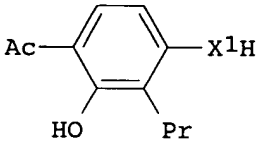

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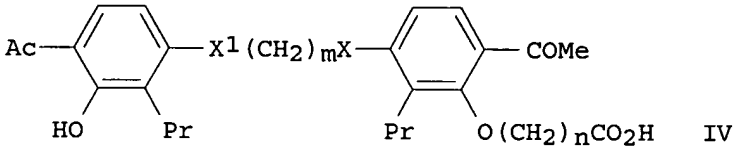


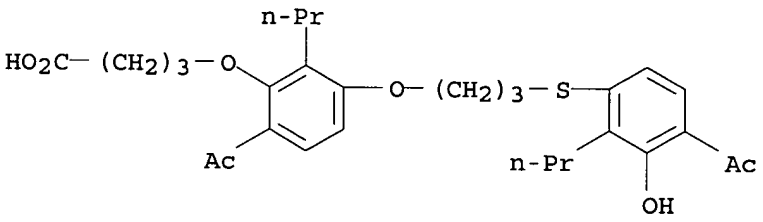
IV

AB The title derivs. IV (m = 2-5; n = 3-8; X1 = S, O; X = O, S, SO, SO2; X1 = X ≠ O), useful as antiallergy agents (no data), are prepared by treating phenoxycarboxylic acids I (Y = halo) with hydroxybenzenes III, which is formed by hydrolysis of hydroxyphenyl carbamates II, in one pot. A mixture of 10 g S-(4-acetyl-3-hydroxy-2-propylphenyl) N,N-dimethylthiocarbamate and KOH in H2O was treated at 95° for 1.5 h, then treated with 12.7 g 4-[6-acetyl-3-hydroxy-3-(3-chloropropoxy)-2-propylphenoxy]butyric acid at 35-40° for 21 h to give 15.2g 4-[6-acetyl-3-(3-(4-acetyl-3-hydroxy-2-propylphentylthio)propoxy)-2-propylphenoxy]butyric acid.

IT **125961-82-2P**
RL: IMF (Industrial manufacture); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of phenoxycarboxylic acid as antiallergy agent from phenoxycarboxylate and hydroxyphenyl carbamate)

RN 125961-82-2 CAPLUS

CN Butanoic acid, 4-[6-acetyl-3-[3-[(4-acetyl-3-hydroxy-2-propylphenyl)thio]propoxy]-2-propylphenoxy]- (9CI) (CA INDEX NAME)



L4 ANSWER 4 OF 7 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 1995:403614 CAPLUS

DOCUMENT NUMBER: 122:290448

TITLE: Preparation of (acetylpropylphenoxy)alkanoic acids as intermediates for antiallergic leukotriene antagonists

INVENTOR(S): Matsumoto, Toyomi; Aizawa, Yasuhiro; Matsukubo, Hiroshi

PATENT ASSIGNEE(S): Kyorin Seiyaku Kk, Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 3 pp.
CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 06345682	A2	19941220	JP 1993-166354	19930611 <--

PRIORITY APPLN. INFO.: JP 1993-166354 19930611
OTHER SOURCE(S): MARPAT 122:290448
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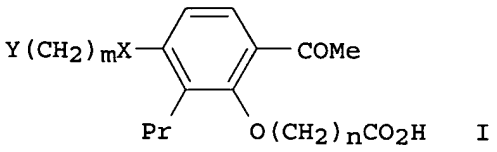
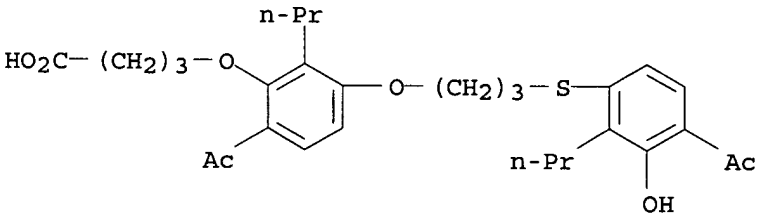


I

AB The title compds. I (m = 2-5; n = 3-8; X = O, S, SO, SO2; Y = halo) are claimed. An aqueous NaOH solution was added dropwise to an EtOH solution of 4-[6-acetyl-3-(3-chloropropoxy)-2-propylphenoxy]butyric acid Et ester (preparation given) at 18-28° and the reaction mixture was stirred at room temperature for 2 h to give 91% 4-[6-acetyl-3-(3-chloropropoxy)-2-propylphenoxy]butyric acid (II). II (21.4 g) and 15.1 g 2-hydroxy-4-mercapto-3-propylacetophenone were dissolved in DMF and the solution was treated with K2CO3 under stirring at room temperature for 3 h to give 24.4 g 4-[6-acetyl-3-[3-(4-acetyl-3-hydroxy-2-propylphenylthio)propoxy]-2-propylphenoxy]butyric acid as a leukotriene antagonist.

IT **125961-82-2P**
RL: IMF (Industrial manufacture); SPN (Synthetic preparation); PREP (Preparation)
(preparation of (acetylpropylphenoxy)alkanoic acids as intermediates for leukotriene antagonists)

RN 125961-82-2 CAPLUS
CN Butanoic acid, 4-[6-acetyl-3-[3-[(4-acetyl-3-hydroxy-2-propylphenyl)thio]propoxy]-2-propylphenoxy]- (9CI) (CA INDEX NAME)



L4 ANSWER 5 OF 7 CAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 1995:39068 CAPLUS
DOCUMENT NUMBER: 123:169347
TITLE: preparation of phenylthiopropoxyphenyloxybutyric acid derivatives as leukotriene antagonists
INVENTOR(S): Oohashi, Mitsuo; Hori, Wataru
PATENT ASSIGNEE(S): Kyorin Seiyaku Kk, Japan
SOURCE: Jpn. Kokai Tokkyo Koho, 13 pp.
CODEN: JKXXAF
DOCUMENT TYPE: Patent
LANGUAGE: Japanese
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 06100526	A2	19940412	JP 1992-273717	19920917 <--

PRIORITY APPLN. INFO.: JP 1992-273717 19920917
OTHER SOURCE(S): MARPAT 123:169347

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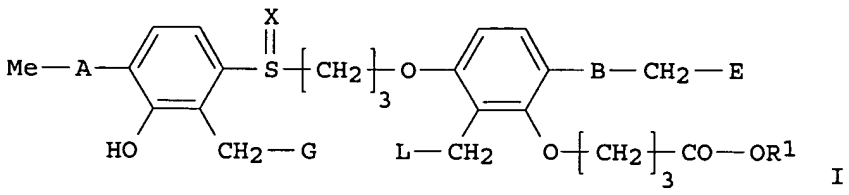


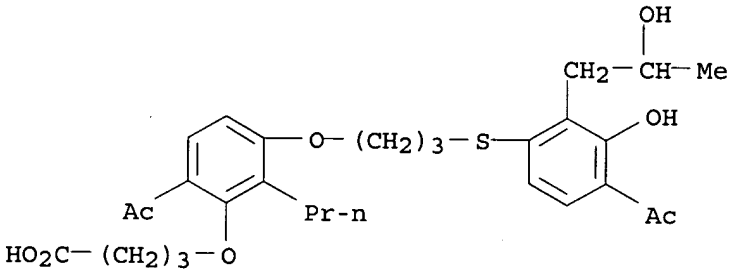
I

AB Title derivs. I (A, B = CO, hydroxymethylene; E = H, OH, acetoxy; G, L = Et, acetyl, 1-hydroxyethyl, 2-hydroxyethyl, hydroxycarbonylmethyl, lower alkoxycarbonylmethyl; X = void, O, O2; R1 = H, lower alkyl; X = O, O2 and B = hydroxymethylene when A = carbonyl, E = H, and G = L = Et) or their alkali salts, acting as strong antagonists for leukotrienes C4, D4, and E4 and useful for antiasthmatics, are prepared Thus, treating 2'-hydroxy-3'-(2-hydroxypropyl)-4'-mercaptoacetophenone (prepared in 6 steps from 3-allyl-2,4-dihydroxyacetophenone) with Et 4-[6-acetyl-3-(3-bromopropoxy)-2-propylphenoxy]butyrate gave I (A = B = CO, E = H, G = 1-hydroxyethyl, L = Et, R1 = Et, X = void).

IT **167211-62-3P 167211-73-6P 167211-79-2P 167211-83-8P 167211-94-1P**
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of phenylthiopropoxyphenyloxybutyric acid derivs. as leukotriene antagonists)

RN 167211-62-3 CAPLUS

CN Butanoic acid, 4-[6-acetyl-3-[3-[[4-acetyl-3-hydroxy-2-(2-hydroxypropyl)phenyl]thio]propoxy]-2-propylphenoxy]- (9CI) (CA INDEX NAME)



RN 167211-73-6 CAPLUS

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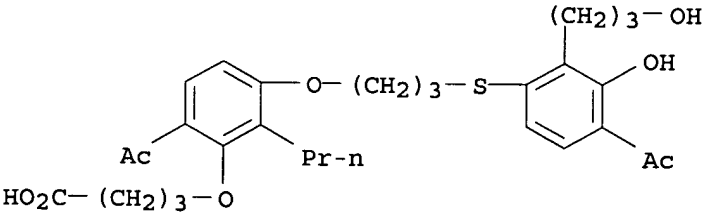
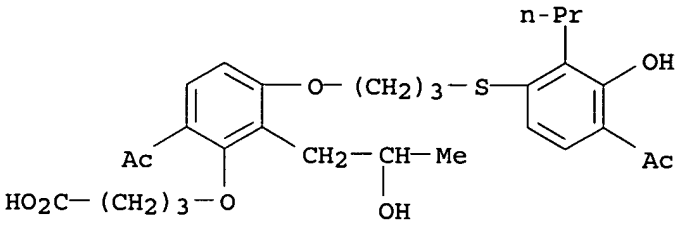


RN 167211-79-2 CAPLUS

CN Butanoic acid, 4-[6-acetyl-3-[3-[(4-acetyl-3-hydroxy-2-propylphenyl)thio]propoxy]-2-(2-hydroxypropyl)phenoxy]- (9CI) (CA INDEX NAME)



RN 167211-83-8 CAPLUS
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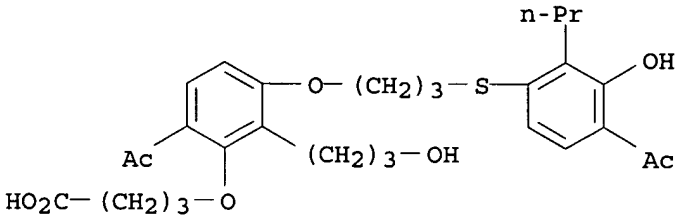
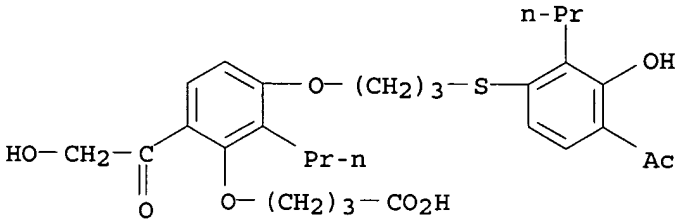


RN 167211-94-1 CAPLUS
CN Butanoic acid, 4-[3-[3-[(4-acetyl-3-hydroxy-2-propylphenyl)thio]propoxy]-6-(hydroxyacetyl)-2-propylphenoxy]- (9CI) (CA INDEX NAME)



L4 ANSWER 6 OF 7 CAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 1990:138760 CAPLUS
DOCUMENT NUMBER: 112:138760
TITLE: Preparation of phenoxyalkylcarboxylic acid derivatives as antiallergic agents
INVENTOR(S): Ohashi, Mitsuo; Awano, Katsuya; Tanaka, Toshio; Kimura, Tetsuya
PATENT ASSIGNEE(S): Kyorin Pharmaceutical Co., Ltd., Japan
SOURCE: Eur. Pat. Appl., 32 pp.
CODEN: EPXXDW
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

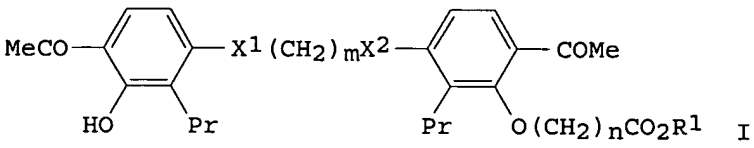
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 332109	A1	19890913	EP 1989-103897	19890306 <--
EP 332109	B1	19911204		
R: BE, CH, DE, ES, FR, GB, IT, LI, NL, SE				
JP 02001459	A2	19900105	JP 1989-38912	19890218 <--
JP 07116125	B4	19951213		
US 4985585	A	19910115	US 1989-313900	19890223 <--
AU 8930884	A1	19890907	AU 1989-30884	19890301 <--
AU 617439	B2	19911128		
CA 1331763	A1	19940830	CA 1989-592555	19890302 <--

HU 50112	A2	19891228	HU 1989-1039	19890303 <--
HU 204030	B	19911128		
HU 208418	B	19931028	HU 1991-2410	19890303 <--
HU 208524	B	19931129	HU 1991-2411	19890303 <--
ES 2045219	T3	19940116	ES 1989-103897	19890306 <--
CN 1036560	A	19891025	CN 1989-101301	19890307 <--
CN 1022407	B	19931013		

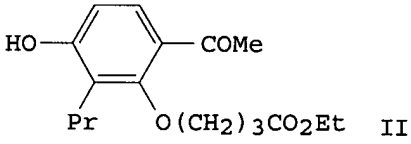
PRIORITY APPLN. INFO.: JP 1988-53374 A 19880307
HU 1989-1039 A3 19890303

OTHER SOURCE(S): MARPAT 112:138760

GI



I



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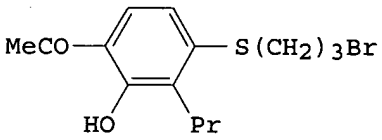


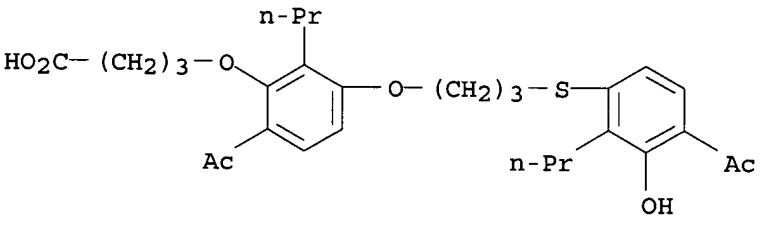
III

AB The title compds. (I; R1 = H, Me, Et; X1, X2 = O, S, SO, SO2; X1 = X2 ≠ O; m = 2-5; n = 3-8), useful as antiallergic agents, are prepared A mixture of phenoxybutyrate II, bromopropyl thioether III, KI, and K2CO3 in Me2CO was refluxed to give 72.4% I (R1 = Et, X1 = S, X2 = O, m = n = 3). I showed 66.7-96.2% inhibition of leukotriene D4-induced bronchoconstriction at 50 mg/kg p.o. in guinea pigs. Addnl. 70 I were also prepared

IT **125961-82-2P 125961-92-4P 125961-93-5P**
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of, as antiallergic agent)

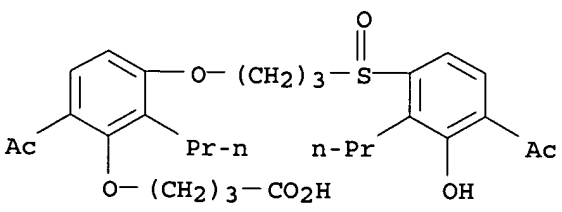
RN 125961-82-2 CAPLUS

CN Butanoic acid, 4-[6-acetyl-3-[3-[(4-acetyl-3-hydroxy-2-propylphenyl)thio]propoxy]-2-propylphenoxy]- (9CI) (CA INDEX NAME)

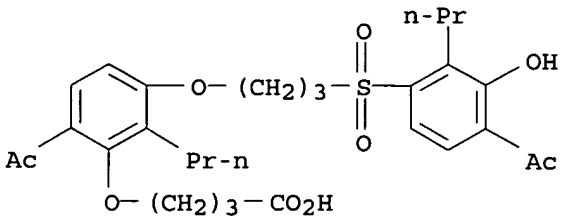


RN 125961-92-4 CAPLUS

CN Butanoic acid, 4-[6-acetyl-3-[3-[(4-acetyl-3-hydroxy-2-propylphenyl)sulfinyl]propoxy]-2-propylphenoxy]- (9CI) (CA INDEX NAME)



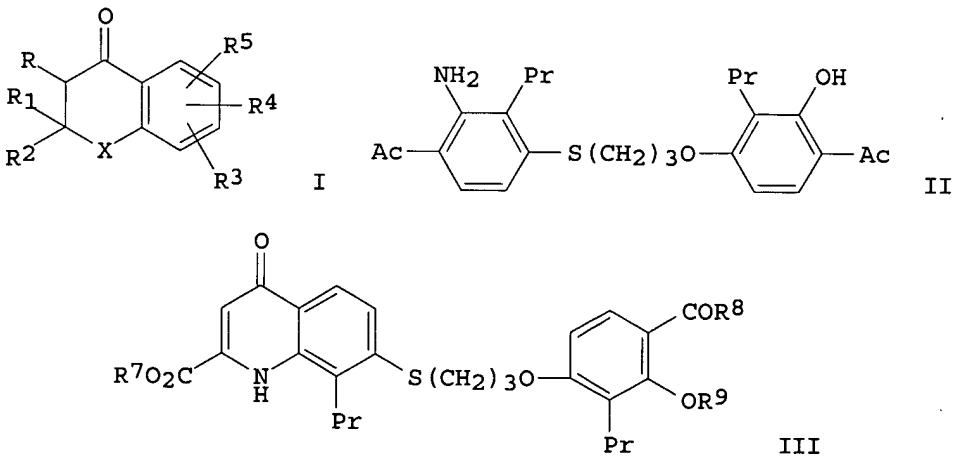
RN 125961-93-5 CAPLUS
CN Butanoic acid, 4-[6-acetyl-3-[3-[(4-acetyl-3-hydroxy-2-propylphenyl)sulfonyl]propoxy]-2-propylphenoxy]- (9CI) (CA INDEX NAME)



L4 ANSWER 7 OF 7 CAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 1983:575604 CAPLUS
DOCUMENT NUMBER: 99:175604
TITLE: Anti-SRS-A carboxylic acid derivatives and pharmaceutical formulations containing them
INVENTOR(S): Bantick, John Raymond
PATENT ASSIGNEE(S): Fisons Ltd., UK
SOURCE: Eur. Pat. Appl., 67 pp.
CODEN: EPXXDW
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

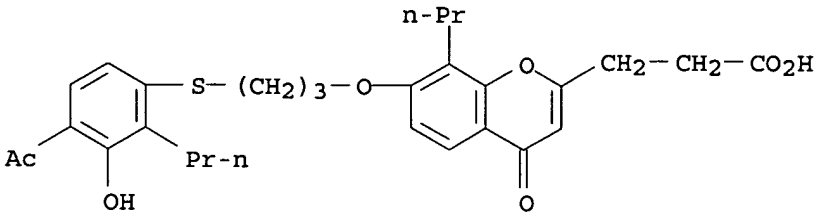
PATENT NO.	KIND	DATE	APPLICATION NO.		DATE	
EP 79637	A1	19830525	EP 1982-201368		19821101	<--
EP 79637	B1	19870128				
R: AT, BE, CH, DE, FR, GB, IT, LI, LU, NL, SE						
US 4474788	A	19841002	US 1982-438163		19821101	<--
AT 25251	E	19870215	AT 1982-201368		19821101	<--
JP 58090557	A2	19830530	JP 1982-196883		19821111	<--
PRIORITY APPLN. INFO.:			GB 1981-34186	A	19811112	
			EP 1982-201368	A	19821101	

GI



AB Anti-allergy (no data) bicyclic compds. I [R, R1 = H, alkyl; RR1 = bond; R2 = CO2H, carboxyalkyl; R3 = substituted OH, SH, NH2; R4, R5 = H, halogen, (un)substituted OH, NH2, alkyl, acyl; X = S, O, NR6 (R6 = H, alkyl)] were prepared Thus, 3,2,4-Pr(HO)2C6H2Ac reacted with 4,2,3-AcPr(H2N)C6H2S(CH2)3Br to give phenol II, which cyclized with EtO2CCO2Et to give quinoline III [R7 = Et, R8R9 = CH:C(CO2Et)]. The latter compound gave III (R7 = H, R8 = Me, R9 = H) on hydrolysis.

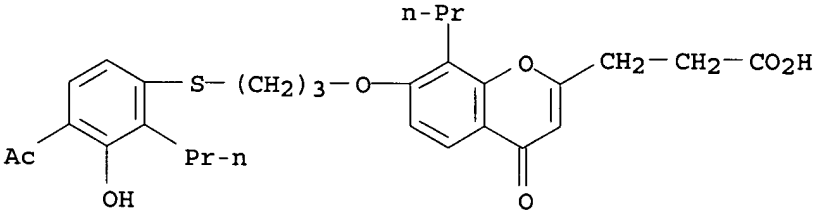
IT^ **87472-35-3P 87472-36-4P**
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)
RN 87472-35-3 CAPLUS
CN 4H-1-Benzopyran-2-propanoic acid, 7-[3-[(4-acetyl-3-hydroxy-2-propylphenyl)thio]propoxy]-4-oxo-8-propyl- (9CI) (CA INDEX NAME)



RN 87472-36-4 CAPLUS
CN L-Lysine, mono[7-[3-[(4-acetyl-3-hydroxy-2-propylphenyl)thio]propoxy]-4-oxo-8-propyl-4H-1-benzopyran-2-propanoate] (9CI) (CA INDEX NAME)

CM 1

CRN 87472-35-3
CMF C29 H34 O7 S



CM 2

CRN 56-87-1
CMF C6 H14 N2 O2

Absolute stereochemistry.